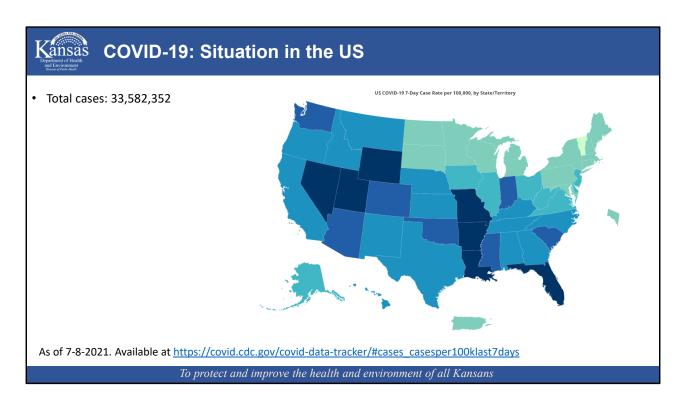


Global Map: https://www.cdc.gov/coronavirus/2019-ncov/locations-confirmed-cases.html.

Last week, we had 182.1 million cases around the world and a bit over 3.9 million deaths.

This week, there are 185,125,237 cases and we have 4,004,006 deaths around the world.



Last week in the US:

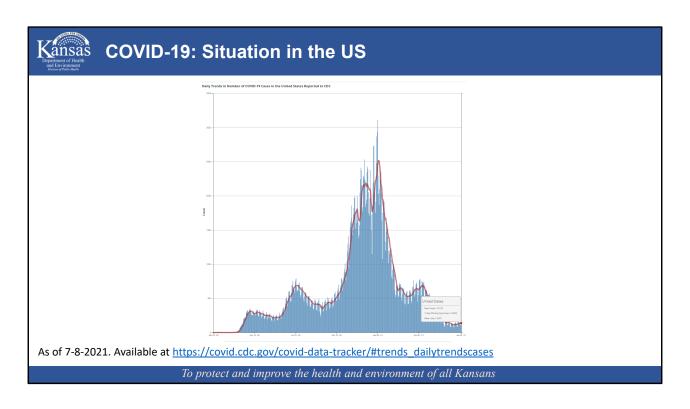
Total cases: 33,485,433 (over 33.4 million)

As of yesterday

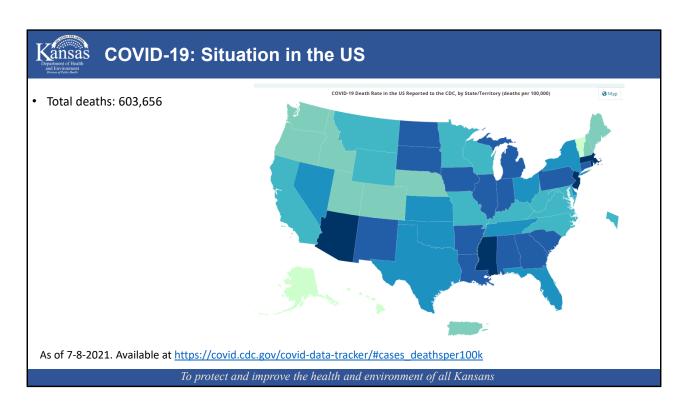
This week:

Total cases: 33,582,352

And you can see from the map of the 7 day case rate that KS is doing okay compared to most of our surrounding states.



In the US, you can see that we are averaging about 13,800 new cases each day according to the 7-day average. That is up from 12,600 from the previous week.



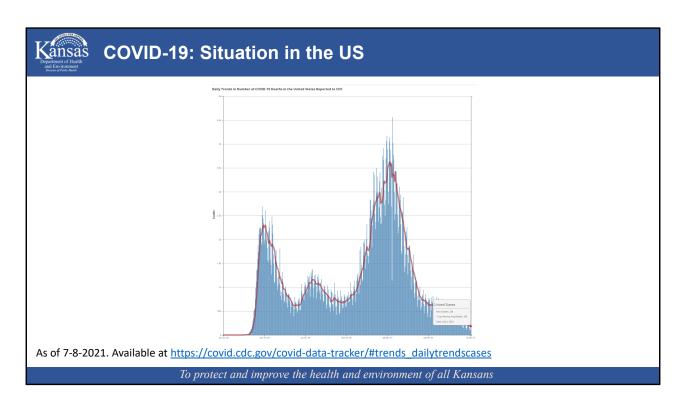
Last week in the US:

Total deaths: 602,133 (a little over 602,000)

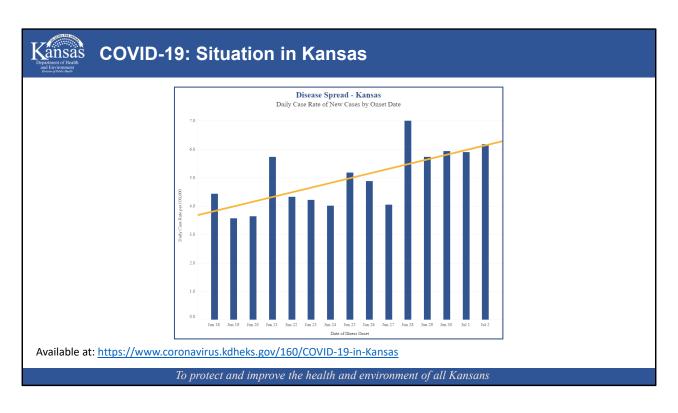
As of yesterday

This week:

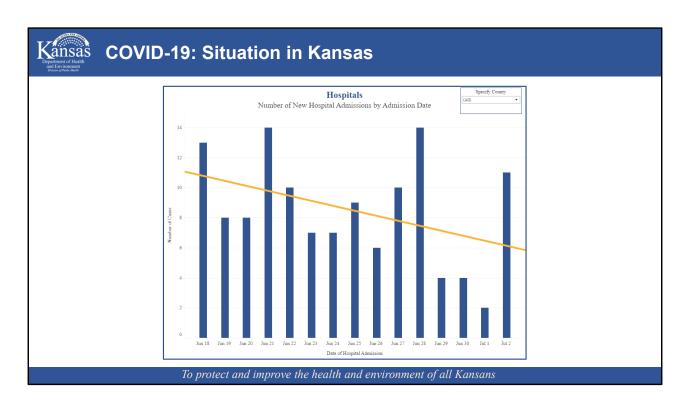
Total deaths: 603,656



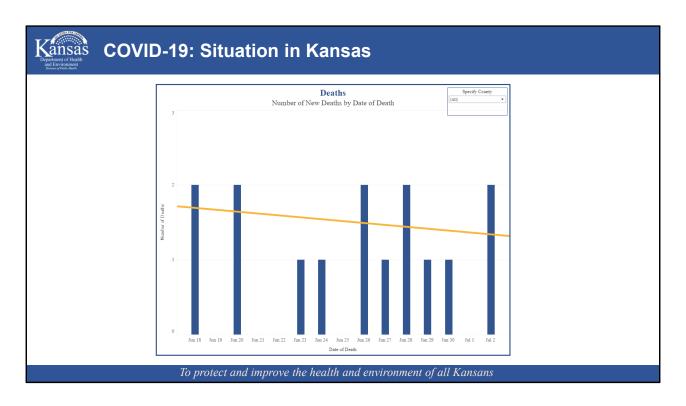
The 7-day moving average daily death trend in the United States is about 183 deaths per day which is down from 280 last week.



Moving on to KS specific data. For our first Disease Spread metric, which is the daily rate of new cases, the trend line last week was slightly increasing last week and it is pretty sharply increasing this week.



For hospitalizations, the trend last week was decreasing last week and that continues this week. As a reminder, this is information gathered during the public health interview of cases and represents hospitalizations at the time of interview. We will take a look at some hospital reported data in a bit to see what that looks like.

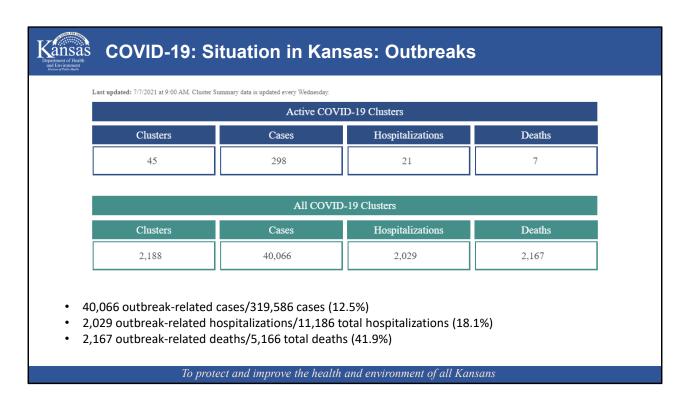


And for deaths, last week the trend was pretty flat and it is slightly decreasing this week.

Kansas COVID-19: Overview							
COVID-19 Cases	Hospitalizations	Statewide Deaths	MIS-C Cases				
319,586	11,186	5,166	17				

As of yesterday, we had 319,586 cases (which is an increase of 1,480 cases since last week) and 5,166 deaths statewide (that's an increase of 10 deaths since last week).

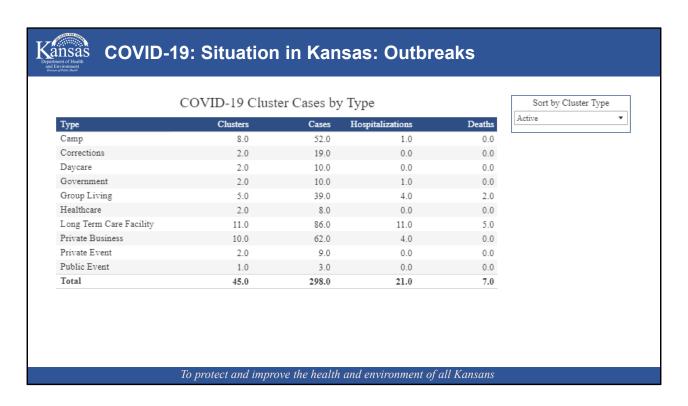
There were 1,071 new cases and 7 new deaths reported between Friday 7/2/2021 and Wednesday 7/7/2021. Monday was a holiday and the dashboard wasn't updated.



Moving on to outbreaks:

As of late Tuesday night, we had 2188 outbreaks across the state; This week we have 45 active clusters which is a big jump from the 26 last week.

Our percentage of outbreak related cases is 12.5%, outbreak-related hospitalizations is about 18.1% and outbreak-related deaths is about 41.9%.

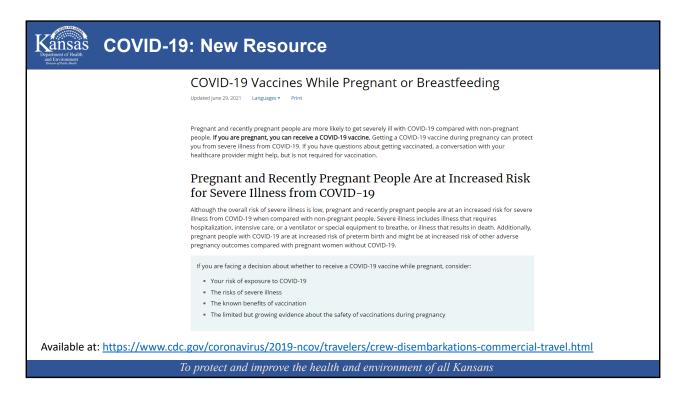


We currently have 8 active outbreaks in camps, which is up from just 1 last week. We have 11 active outbreaks in LTCFs (which is up slightly from 9 last week). We also have 5 outbreaks in group living settings and 10 in private businesses.

Don't forget, if you are interested in seeing the list of named locations with 5 or more cases within the last 14 days, you can go to the dashboard.

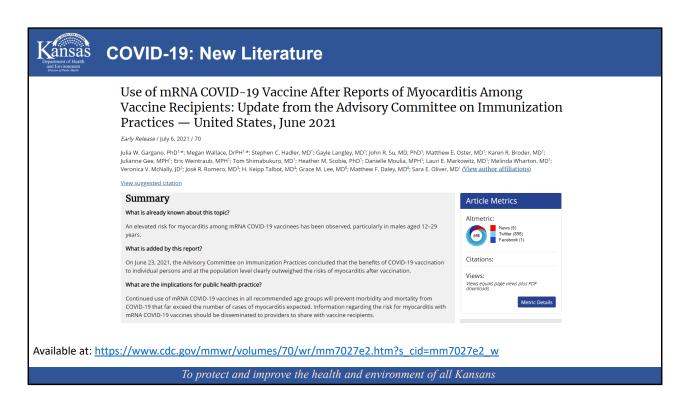
Kansas COVID-19): New F	Resour	ce			
	This table is updated sev	eral times a week, as	needed. Last updated July 7, 2021	with EDC data subn	nitted July 6, 2021	
	Parent Company	Cruise Line	No Sail Order Response Plan Status	Ship Name	Ship Status	
	Bahamas Paradise Cruise Line	Bahamas Paradise Cruise Line	Complete and accurate with signed acknowledgement	Grand Classica	Red	
	Carnival Corporation	Carnival Cruise Lines, Inc.	Complete and accurate with signed acknowledgement	Carnival Breeze	Orange	
				Carnival Conquest	Green	
				Carnival Ecstasy	Green	
				Carnival Elation	Green	
				Carnival Freedom	Orange	
				Carnival Horizon	Orange	
				Carnival Liberty	Orange	
				Carnival Mardi Gras	Orange	
Available at: https://www.cdc.gov/coronavirus/2019-ncov/travelers/crew-disembarkations-commercial-travel.html						
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CDC implemented the COVID-19 color-coding system for cruise ships to mitigate transmission of COVID-19 onboard. Onboard <u>preventative measures</u> are recommended or required based on ship color status.



The June 29, 2021 update provides information on benefits and side effects related to COVID-19 vaccines, and available data on vaccination among people who are pregnant, those who are breastfeeding, and those would like to have a baby.

There is currently no evidence that any vaccines, including COVID-19 vaccines, cause female or male fertility problems—problems getting pregnant. CDC does not recommend routine pregnancy testing before COVID-19 vaccination. If you are trying to become pregnant, you do not need to avoid pregnancy after receiving a COVID-19 vaccine.



In May 2021, FDA expanded the EUA for the Pfizer-BioNTech COVID-19 vaccine to include adolescents aged 12–15 years; ACIP recommends that all persons aged ≥12 years receive a COVID-19 vaccine. After reports of myocarditis and pericarditis in mRNA vaccine recipients,¶ which predominantly occurred in young males after the second dose, an ACIP meeting was rapidly convened to review reported cases of myocarditis and pericarditis and discuss the benefits and risks of mRNA COVID-19 vaccination in the United States.

On June 23, 2021, the Advisory Committee on Immunization Practices concluded that the benefits of COVID-19 vaccination to individual persons and at the population level clearly outweighed the risks of myocarditis after vaccination. The EUA has been modified to include information on myocarditis after receipt of mRNA COVID-19 vaccines. The EUA fact sheets should be provided before vaccination; in addition, CDC has developed patient and provider education materials about the possibility of myocarditis and symptoms of concern, to ensure prompt recognition and management of myocarditis.

Kansas COVID-19:	New Literature
	Annals of Epidemiology Available online 20 June 2021 In Press, Journal Pre-proof ⊕
	Trends in the distribution of COVID-19 deaths by age and race/ethnicity — United States, April 4— December 26, 2020
	Lauren M. Rossen PhD ¹ 名思, Jeremy A.W. Gold MD ² , Farida B. Ahmad MPH ¹ , Paul D. Sutton PhD ¹ , Amy M. Branum PhD ¹ Show more
	+ Add to Mendeley <\$ Share 55 Cite https://doi.org/10.1016/j.annepidem.2021.06.003 Get rights and content
	Key words Mortality, Coronavirus 2019, SARS-CoV-2; Disparities; National Vital Statistics System
	Introduction
	The coronavirus disease 2019 (COVID-19) pandemic has disproportionately affected racial and ethnic minority groups.1, 2, 3, 4, 5 COVID-19 infection and mortality rates are higher among Hispanic/Latino, non-Hispanic Black, and non-Hispanic
Available at: https://www.scienced	direct.com/science/article/pii/S104727972100154X?via%3Dihub
То д	protect and improve the health and environment of all Kansans

Key findings:

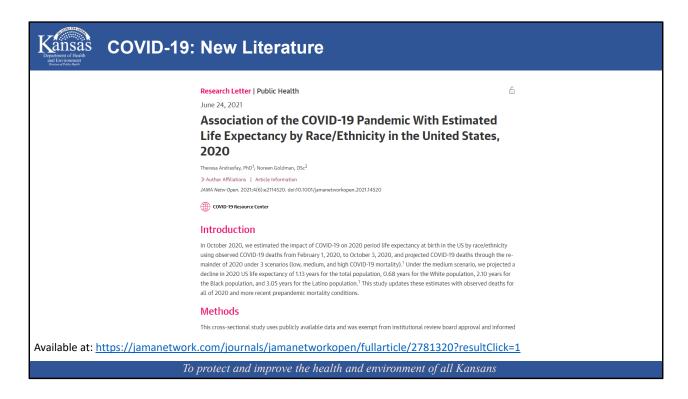
The largest percentage of COVID-19 deaths from April through October occurred in older persons aged ≥ 85 years among both the non-Hispanic White and non-Hispanic Asian populations (Figure).

Compared with the White population, the age distributions of COVID-19 deaths among non-Hispanic NHOPI and other racial/ethnic groups, were younger (Figure).

The percentage of COVID-19 deaths declined in all age groups 25 and older for non-Hispanic Black persons since April and, since late July, for Hispanic persons.

Methods: Using data from the National Vital Statistics System, the percentage distribution of 363,087 US COVID-19 deaths from April 4 through December 26, 2020 was calculated by age group (<25, 25–44, 45–64, 65–74, 75–84, and ≥85 years) within race/ethnicity groups (Hispanic, non-Hispanic White, non-Hispanic Black, non-Hispanic Asian, non-Hispanic Al/AN, non-Hispanic Native Hawaiian or other Pacific Islander [NHOPI], other). *Limitations*: COVID-19 deaths may be underestimated by race/ethnicity due to differential reporting on death certifications; small number of deaths in some race/ethnic groups.

Implications: Prevention efforts need to focus on highly impacted older age groups and also on Hispanic/Latino, non-Hispanic Black, non-Hispanic AI/AN, and non-Hispanic NHOPI populations, where a majority of COVID-19 deaths were among persons aged <75 years.

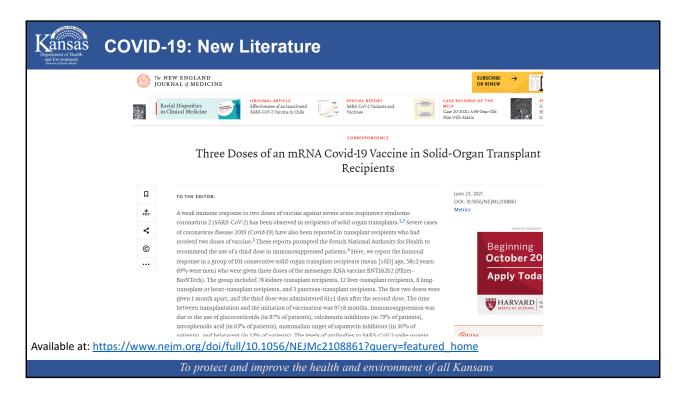


Key findings:

Updated life expectancy estimates show that COVID-19 reduced overall US life expectancy from 78.74 to 77.43 years (Figure).

Compared with the White population (0.94 years), reductions were 3.2 times larger for the Latino population (3.03 years) and 2 times larger for the Black population (1.90 years) (Figure).

Methods: Cross-sectional study using 380,868 COVID-19 deaths in 2020 to estimate changes in life expectancy associated with COVID-19 for the total US population, non-Latino White, non-Latino Black, and Latino populations. *Limitations*: COVID-19 deaths may be underestimated by race/ethnicity due to differential reporting on death certificates. **Implications**: Disproportionate declines in life expectancy among Latino and Black populations may stem from social and economic inequities that are associated with higher exposure to infection, poorer care once infected and thus higher fatality among those infected.



The next few pieces of literature are being presented to you for your information. There are a lot of questions about booster doses, booster doses for certain susceptible populations, what are the effects of mixing and matching vaccine, etc. I have found a few research projects looking at these questions and am presenting articles today. This does not mean that KDHE is making any new recommendation about how vaccines should be administered. But since these questions come up a lot, I am including some information on scientific studies here. I will continue to bring these to you as I find them.

Key findings:

Among solid-organ transplant recipients, the prevalence of anti-SARS-CoV-2 antibodies was:

0% (95% CI 0 to 4) of 101 patients before the 1st dose of BNT162b2 (Pfizer/BioNTech) (Figure).

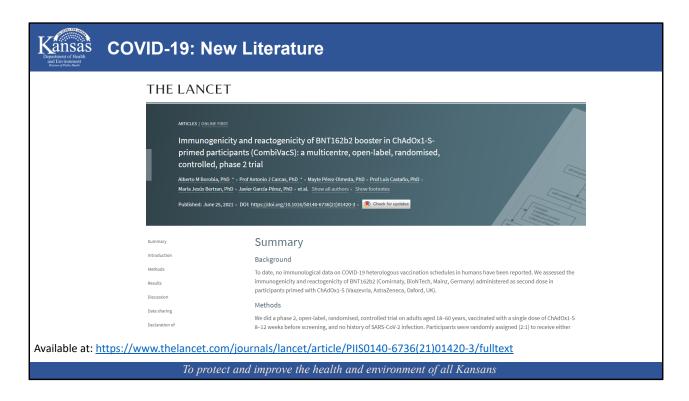
4% (95% CI 1 to 10) of 101 patients before the 2nd dose (Figure).

40% (95% CI 31 to 51) of 99 patients before the 3rd dose (Figure).

68% (95% CI 58 to 77) of 99 patients 4 weeks after the 3rd dose (Figure).

No serious adverse events were reported after the 3rd dose, and no acute rejection episodes occurred. **Methods:** In an anonymous retrospective study of solid-organ transplant recipients in France, 99 patients received a 3rd dose of BNT162b2 vaccine 61±1 days after the 2nd dose. Prevalence of anti-SARS-CoV-2 antibodies was assessed before the 3rd dose and 1 month afterward. Any adverse events, including rejection episodes, were recorded. *Limitations*: Small sample size, unknown how antibody response correlates with protection from infection.

Implications: Immunogenicity to BNT162b2 is reduced among solid-organ transplant patients but increases with additional doses. Transplant recipients and their close contacts should be vaccinated against COVID-19, and transplant recipients should continue to take precautions (masking, distancing, avoiding crowds and poorly ventilated indoor spaces) to protect themselves against COVID-19, even after vaccination.



Key findings:

By day 14 post-heterologous COVID-19 vaccination:

SARS-CoV-2 receptor binding domain antibodies (anti-RBD) titers increased from 71.46 BAU/mL to 7,756.68 BAU/mL.

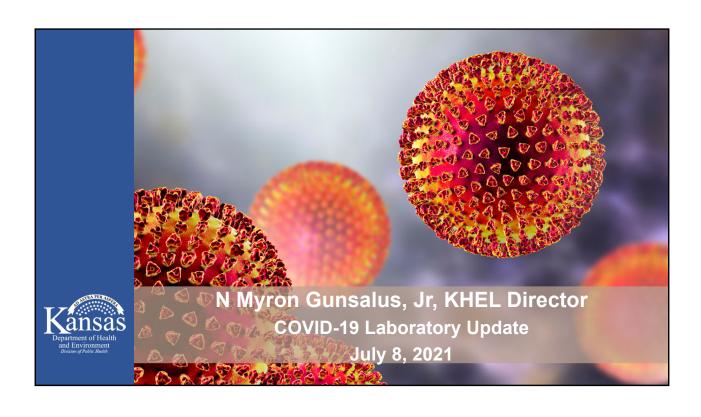
Trimeric spike protein antibody titers increased from 98.40 BAU/mL to 3,684.87 BAU/mL. Neutralizing antibody capacity was high ($NT_{50}>1:300$ and <1:1,000) or very high ($NT_{50}>1:1,000$).

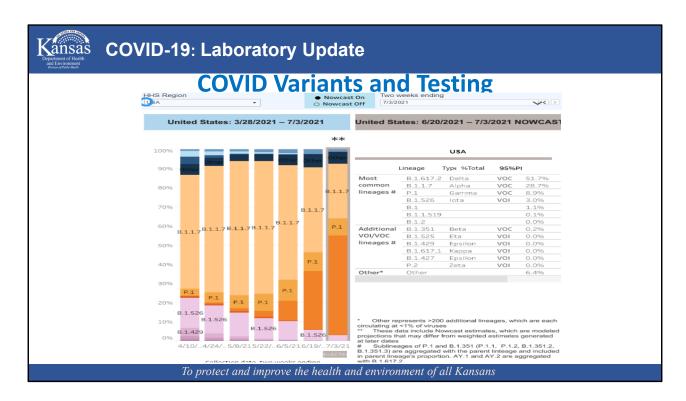
All humoral immune responses were significantly higher than those of the controls (p<0.0001) (Figure).

No serious adverse events were reported.

Methods: Phase 2, open-label, randomized, controlled trial conducted in 5 Spanish hospitals. Adults aged 18–60 years who received a prime ChAdOx1 (AstraZeneca/Oxford) dose 8–12 weeks prior to enrollment were randomly assigned (2:1) to receive either a single dose of BNT162b2 (Pfizer/BioNTech, n = 441) or no vaccine (control group, n = 222). Immune response and antibody neutralization capacity (n = 198) were determined up to 14 days after BNT162b2 dose. 7-day reactogenicity was also determined. *Limitations*: No control group completing the homologous ChAdOx1 regimen.

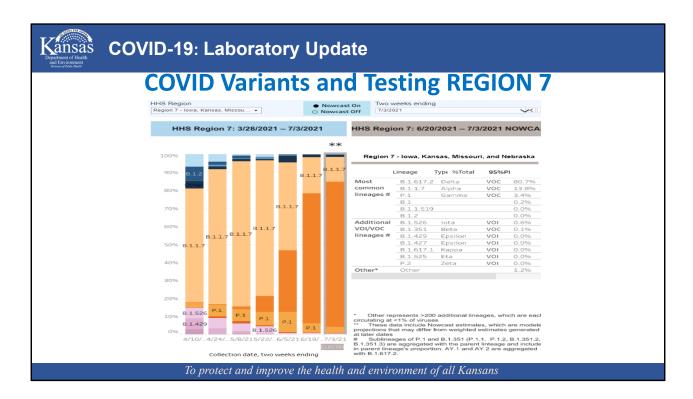
Implications: The use of an mRNA vaccine as a 2nd dose in a heterologous scheme may increase the immune response obtained after an initial dose of ChAdOx1, without serious adverse events.





 $tracker/?CDC_AA_refVal=https\%3A\%2F\%2Fwww.cdc.gov\%2Fcoronavirus\%2F2019-ncov\%2Fcases-updates\%2Fvariant-surveillance\%2Fgenomic-surveillance-dashboard.html \#variant-proportions$

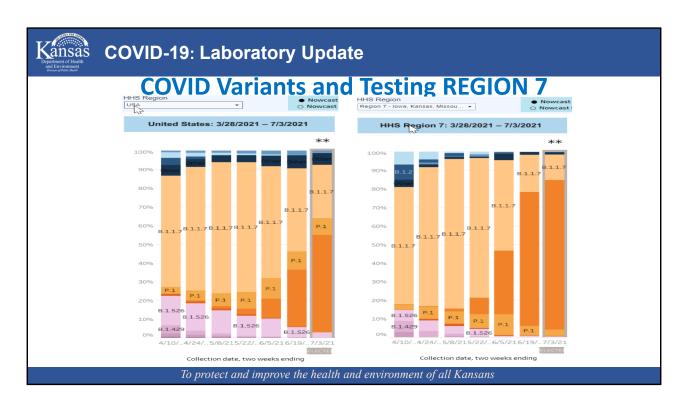
This is the latest with prediction/now cast on to predict out. Significance is that P.1 and B.1.617.2 are growing significantly.



 $tracker/?CDC_AA_refVal=https\%3A\%2F\%2Fwww.cdc.gov\%2Fcoronavirus\%2F2019-ncov\%2Fcases-updates\%2Fvariant-surveillance\%2Fgenomic-surveillance-dashboard.html \#variant-proportions$

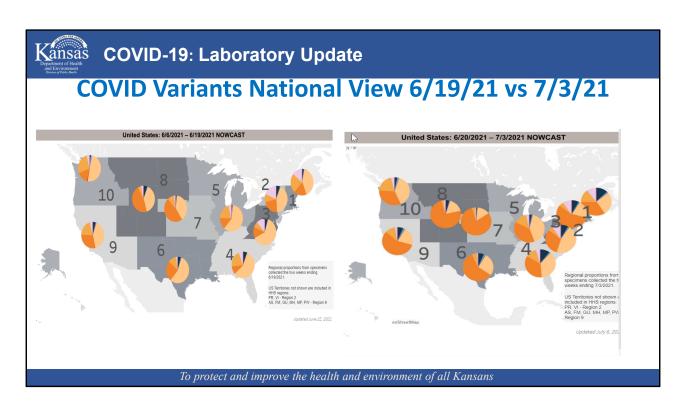
CDC now has a new modeling program included called "Nowcast". If you turn it on, as shown in this slide, it predicts the next 2 weeks of proportional data associated with the variants. You can use it for regional or US wide evaluations.

Keep in mind these are not absolute numbers but are proportional numbers and only relate to those sequences performed by CDC

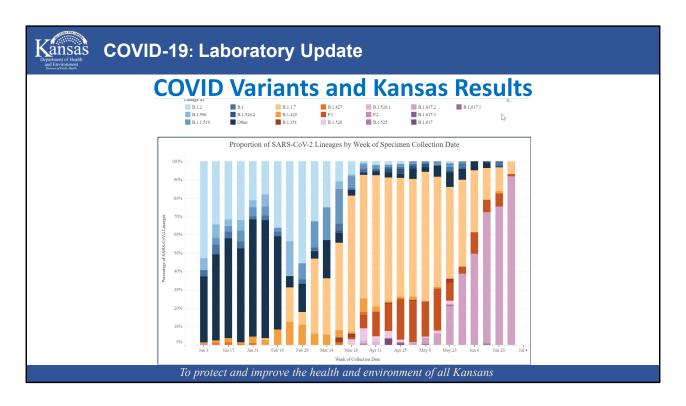


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This is comparing USA vs Region 7

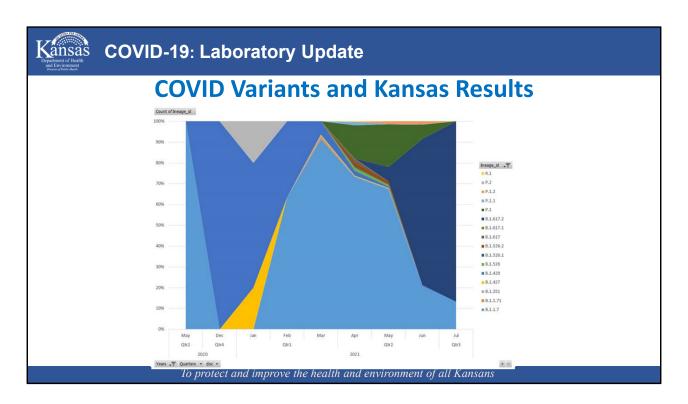


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https://www.coronavirus.kdheks.gov/160/COVID-19-in-Kansas

The purple is the Delta, B.1.617.2 variant



Medium blue in Dec/January was mostly B.1.429 or California strain.

Large lighter blue in the middle is the B.1.1.7 (Alpha) strain

The Dark Blue that is currently ~80% of the total is B.1.617.2 or Delta Variant.

This graph is only VOC. But to give you a better idea we have only sequenced 7 non-VOCs since June 25th! 7 out of 377 in that time period are non-VOCs. So this graph is pretty accurate to the overall.

Y axis is percentage. B.1.1.7 (alpha) is dropping rapidly and being taken over by B.1.617.2 (delta).

If you are a lab that is running PCR for COVID detection, we are looking for ways to increase our statewide sequencing efforts



KHEL is looking to partner with labs to increase sequencing



Criteria

KHEL is asking labs to send any positive samples

However, KHEL is particularly interested in the following cases if all samples are not available

• Examples: Areas with high transmission (different age groups, geographic locations, severity), cases in areas with a significant increase of cases over a few weeks (not explained by relaxing public health measures), children in areas with increased incidence of pediatric disease, clusters of cases in people aged <60 without underlying conditions, cases in fully-vaccinated people or when reinfection is suspected

Samples with a CT value <30 are preferred, but higher CT values are OK if samples are of interest



Next steps

- To sign up for the program, Contact KDHE.KHELINFO@KS.gov and include Subject Line: ATTENTION SEQUENCING
 - Samples can be submitted through a form or lab online
 - For regular surveillance tell us how many PCR positives you typically have per week and could send.
- If there is a known case of reinfection or potential vaccine break through or "S-Deletion", then contact KHEL for sequencing.
- You should not report PCR mutation screening to anyone as an identified variant. Variants are only identified after confirmation whole genome sequencing
 - Send us extract if possible or a second sample in VTM

We will be providing guidance over next week to send all positive samples with a CT of less than 30 to KHEL for sequencing. We may or may not be able to do all of them, but this way we will start to get a better cross section of what is going on in the state.

We are currently working on a system that will also allow us to report back with the understanding that any sequencing data should not be used for patient care or diagnostic evaluation but can be used for public health response.

Please note: CMS has stated so far that sequencing results cannot be reported back to providers but only to Public Health. We will be working with Public Health staff regarding results of sequences but can only provide aggregate data back to any given laboratory



COVID-19: Laboratory Update---Courier Update

- It is live now. All counties should have a pickup happening 5 days per week.
- We do not want to delete any pickup site for at least the first month of full service.

Any questions regarding the courier service can be directed to:

Chad Yamashita

E-mail: Chad.Yamashita@ks.gov

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Kansas COVID-19: Laboratory Update

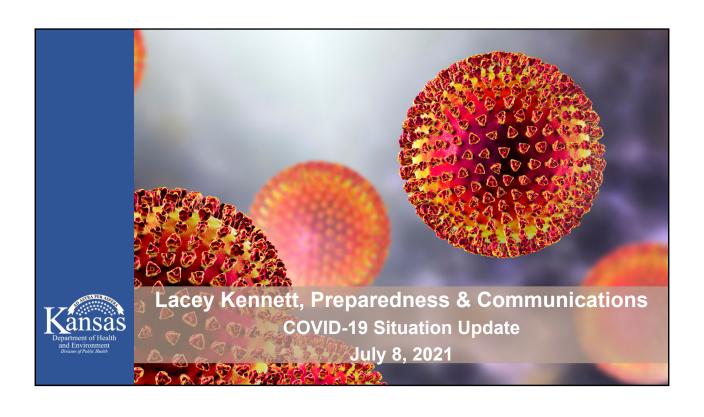
Updates and Reminders

- CLIA Certification Questions: KDHE.CLIA2@ks.gov
 - REMINDER: If you add or change anything on your test menu, you must notify the CLIA office ASAP. (even if High Complexity lab adding antigen.)
- Kits and some instruments available
- Mobile Labs and Collection Vans available.

To protect and improve the health and environment of all Kansans

We were notified by the CLIA certification office that there are a number of labs that have either Certificates of Waiver(COW) or other more complex certifications that brought on the antigen testing such as BinaxNow without notifying the CLIA office of the change in their test menu. All labs (including COWs) are required to notify CLIA of any addition or subtractions of tests or technologies offered in their test menu.

Adding COVID testing with the CLIA office does not issue a new certificate or put the name of the test on your certificate. It is a procedural requirement.





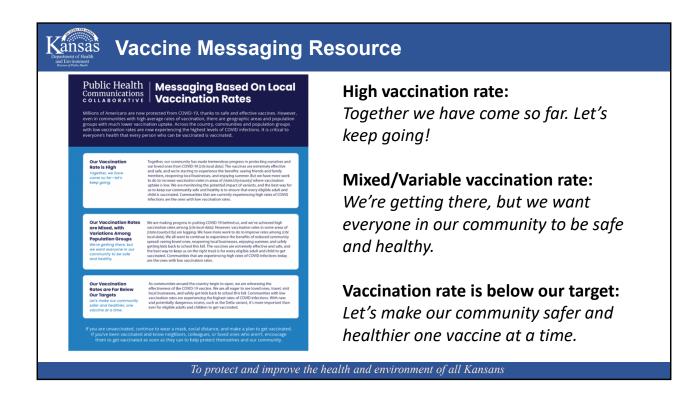
On Wednesday, July 21, the Public Health Communications Collaborative will present a webinar called "COVID-19 and Return to In-Person Learning: Communicating with Constituents." This webinar will take place at 12:00 noon Central time. The webinar will focus on effective messaging to communicate with families, teachers, and school officials in preparation for a safe and comfortable return to the classroom. Representing different local scenarios and constituencies, our three panelists will share insights and answer your questions. To register for this free webinar, click the link on your screen or visit https://trustforamericashealth.webex.com/mw3300/mywebex/default.do?nomenu=true& siteurl=trustforamericashealth&service=6&rnd=0.8075004991391943&main_url=https%3A%2F%2Ftrustforamericashealth.webex.com%2Fec3300%2Feventcenter%2Fevent%2Fevent Action.do%3FtheAction%3Ddetail%26%26%26EMK%3D4832534b0000000509387d18c95c7c8e53147e7da5fc9c15d9890a6e1fd2acb48f26e23192c0453b%26siteurl%3Dtrustforamericashealth%26confViewID%3D198392026881888944%26SourceId%3DphccTwitter%26encryptTicket%3DSDJTSwAAAAX6Pgem4336hiNWHDwQiGvAtAT5wEuLBHq_p4N1a-rVbw2%26.



The latest Public Health On Call podcast from Johns Hopkins University is now available.

"COVID-19 Vaccines and Kids" - Youths 12 and older have been eligible for COVID-19 vaccines since March, but clinical trials are still ongoing for kids under 12. Dr. Kawsar Talaat, who led one of Pfizer's COVID-19 vaccine trials in adults, and Dr. Odis Johnson, executive director of the Johns Hopkins Center for Safe and Healthy Schools, talk about whether schools may require vaccines, the ethics of immunizing children when so many high-risk adults around the world don't have access to shots, risk factors for serious disease among children, and what is known currently about vaccine hesitancy among parents. Link: https://johnshopkinssph.libsyn.com/342-covid-19-vaccines-and-children

You can listen to the podcast by clicking the link or wherever you listen to podcasts.



Also from the Public Health Communications Collaborative, there is a new tool that can help you message vaccines to your community members based on local vaccination rates. You can see a screenshot of the messaging advice here, but a full-size PDF is available for download. Vaccine messaging for your community will vary depending on whether your community has a high vaccination rate, a low vaccination rate, or somewhere in between and this document can help. I've put a short summary of each category on this slide, but the document has so much more information, so I encourage you to download it. You can access it here: https://publichealthcollaborative.org/resources/graphic-messaging-based-on-local-vaccination-rates/



Conspiracy Theorists Claim Moderna Vaccine Was Developed Pre-Pandemic

July 1, 2021

False claims that the Moderna COVID-19 vaccine was developed before the pandemic have been circulating online for the past two weeks. These claims are based on an agreement between Moderna, the University of North Carolina at Chapel Hill, and the National Institutes of Allergy and Infectious Diseases (NIAID) that outlines the terms for transferring research materials related to coronavirus RNA vaccine between the institutions. Because the agreement was signed prior to WHO notification of COVID-19 in China, it is being used as proof of a "plandemic," the conspiracy theory that the COVID-19 pandemic was planned. According to a spokesperson from the NIAID, the agreement pertained to vaccine candidates against another coronavirus, Middle East Respiratory Syndrome coronavirus (MERS-CoV), and not SARS-CoV-2. NIAID and Moderna have been collaborating on coronavirus vaccine research since 2017, well before the identification of SARS-CoV-2.

Recommendation: Ignore Read More +

Recommendations are provided, organized into three categories:

- Ignore: Focus on current communications priorities.
- Passive Response: Be prepared to address if directly asked, and in certain cases consider updating FAQ's and info sheets addressing common myths and misperceptions.
 Otherwise, continue to focus on current communications priorities.
- Direct Response: Directly address this misinformation.

Bookmark THIS LINK

To protect and improve the health and environment of all Kansans

This is another great resource from the Public Health Communications Collaborative that you might find helpful. They have a website dedicated to tracking COVID-19 misinformation that is found online and in social media. The website summarizes the misinformation, discusses its origins, and gives recommendations on whether it is best to ignore the misinformation, be prepared for a passive response, or directly address the misinformation. There is also a section called "Tough Q&A" that can help you formulate an answer to some of the tough questions you might get about COVID-19. All of this is available on their website: https://publichealthcollaborative.org/misinformation-alerts/



This resource would be a great tool to share on social media or with technology-savvy friends and family.

Vira ("Vaccine Information Resource Assistant") is a chatbot developed by the International Vaccine Access Center at the Johns Hopkins Bloomberg School of Public Health with experts from the Bloomberg School's Department of International Health and Johns Hopkins Whiting School of Engineering.

Vira's knowledge base from Johns Hopkins vaccine scientists was developed with support from the COVID-19 Training Initiative, with funding from Bloomberg Philanthropies. VaxChat is powered by IBM Technology. You can visit VaxChat.org and as Vira questions about the COVID vaccine, and she'll provide you science-based answers. Vira is also a useful technology for the Bloomberg School of Public Health, as it monitors the questions asked and helps scientists develop answers to the questions that people have the most.

You can see on the right hand side of this screen I asked Vira a simple question and she provided me with an answer and an opportunity to give feedback about its answer. The questions and answers she gives are publicly available on a companion app called the COVID-19 Vaccine Resource Hub. There are also browsable FAQs on the VaxChat.org website.



Rural telehealth adoption is growing significantly – particularly by connecting patients in their homes to a provider. While the growth is compelling, there is still untapped potential for rural hospitals to utilize telemedicine to drive down costs and improve the delivery of care within the acute care setting. This webinar will discuss ways to leverage telehospitalists to stabilize staffing, reduce recruiting costs, support advanced practice providers, and positively impact sub-specialist care. The webinar is hosted by the National Rural Health Association on Wednesday, July 14 at noon CT.

To register: https://register.gotowebinar.com/register/5710471284127098126

